



✓ Site: Philadelphia

# FDAMA STAKEHOLDER MEETING

APRIL 28, 1999

## Talking with Stakeholders About FDA Modernization

\*\*Your question/comments will become part of Docket Number: 99N-0386

Fax to: 1-888-361-4011 (on April 28 only)

Title (required) First Name (required) Last Name (required)  
☒ Dr. ☐ Mr. ROBERT DODEMAIDE  
☐ Mrs. ☐ Ms.  
 Organization HOECHST ROUSSEL VET

Stakeholder Group ✓ stakeholder group you represent

☐ Consumer ☐ Consumer Group ☐ Health Professional ☐ Industry ☐ Association ☐ Other

Center ✓ the center/product area your comments address

☐ Center for Biologics ☐ Center for Drug Evaluation and Research  
☐ Center for Devices and Radiological Health ☐ Center for Food Safety and Applied Nutrition  
☒ Center for Veterinary Medicine ☐ Office of Regulatory Affairs  
☐ FDA General

### Questions to Stakeholders

Please check the box next to the stakeholder question/s from the March 22, 1999, Federal Register notice which your question/comment addresses.

- ☒ 1. What actions do you propose the Agency take to expand FDA's capability to incorporate state-of-the-art science into its risk-based decision-making?
- ☐ 2. What actions do you propose to facilitate the exchange and integration of scientific information to better enable FDA to meet its public health responsibilities throughout a product's life cycle?
- ☐ 3. What actions do you propose for educating the public about the concept of balancing risks against benefits in public health decision-making?
- ☐ 4. What actions do you propose to enable FDA and its product centers to focus resources on areas of greatest risk to the public health?
- ☐ 5. What additional actions do you propose for enhancing communication processes that allow for ongoing feedback and/or evaluation of our modernization efforts?
- ☐ 6. Additional Comments on FDA Modernization Efforts.

### YOUR COMMENT/QUESTION

While funds were available, CVN1 provided "externships" to its staff. These were greatly appreciated by both CVN1 ("no idea this is how industry works") and by industry ("good opportunity to show our capabilities to FDA").

I encourage the FDA to build upon this modest beginning with 1- or 2-week duration externships.

99N-0386

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